CDC Update

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CLIAC Meeting April 15, 2015 Atlanta, Georgia



Division of Laboratory Systems

- Dual Use Research of Concern
- Specimen management (CASPIR)
- Instructional Design
- Training Distribution
- Web Broadcast
- CDC TRAIN Assistance

Support Services for CDC Research and Education

- Clinical laboratory and public health/healthcare integration
- Regulatory and voluntary standards development
- Training- public health and clinical workforce

- Measurement of training effectiveness
- Measurement of impact of laboratory practice guidelines and laboratory based continuing education programs

Evaluation

Innovation

- Multimedia enhanced distance learning
- Clinical decision support
- Passive surveillance with CMS data
- Collaboratives for genetic testing (GeT-RM) and Next-Gen sequencing

DLS Role in Ebola Response

- Advice on clinical laboratory issues, especially safety
 - Support laboratories, clinical and public health, both at home and abroad
 - Coordinate with CDC Emergency Operations Center laboratory teams
 - Videos demonstrating biosafety procedures
 - Amer. Biol. Safety Assoc. certified staff supports triage/ response to inquiries from laboratories and field epidemiologists

DLS and CDC Laboratory Safety Efforts

- Supporting Laboratory Safety workgroup recommendations
 - Creating improved laboratory training in
 - Safety and quality management
 - Competency-based curriculum
 - Laboratory Leadership Service fellowship CSELS/DLS & DSEPD
- External accreditation for all CDC laboratories
 - CLIA for clinical labs; ISO17025 for research labs
- CASPIR, "CDC and ATSDR Specimen Packaging, Inventory and Repository"
 - Advancing improved IT accession systems, with bar coding
 - Revising specimen collection

Promoting Good Laboratory Practices for Waived Testing - Update to 2014 CLIAC Recommendation

- Development of a voluntary selfassessment checklisttype tool based on recommended practices in the "Ready? Set? Test!" booklet
- Updates being made to "Ready? Set? Test!" and "To Test or Not To Test?" booklets to address offlabel use of waived tests

Tips, Reminders, and Resources

READY?

- ☐ Clean work surfaces before and after testing.
- ☐ Perform testing in a well lit area.
- Check and record temperatures of the testing and reagent storage areas.
- Check inventory regularly to ensure you will have enough reagents and supplies on hand for testing.
- Check and record expiration dates of reagents/kits, and discard any reagents or tests that have expired.
- Check that all kit reagents came from the same kit lo Do not mix reagents.
- ☐ Inspect reagents for damage, discoloration, or contamination, and discard if found.
- ☐ Prepare reagents according to manufacturer's instructions.
- $\hfill \square$ Allow time for refrigerated reagents/samples to come to room temperature prior to testing.
- ☐ Inspect equipment and electrical connections to be sure they are working.
- ☐ Perform calibration checks, as needed, following the manufacturer's instructions.
- ☐ File the old manufacturer's instructions and replace with the new copy if there are changes.
- Communicate all changes in the manufacturer's instructions to other testing personnel and to the person who directs or supervises testing.
- ☐ Treat and test quality control (QC) samples the same as patient samples.
- ☐ Perform QC as recommended in the manufacturer's instructions.

SET?

- ☐ Check patient identification and test orders.
- ☐ Discuss pretest instructions and counseling needs with the patient.
- ☐ Wear appropriate personal protective equipment (PPE) such as gloves.
- Collect and label a good sample for testing.
- ☐ Clean hands and change gloves between patients.
- ☐ Use the proper biohazard containers to dispose of waste and sharps.

TEST!

- ☐ Do not test samples that are improperly collected or handled.
- ☐ Have the manufacturer's instructions or a quick reference guide at the work station.
- ☐ Follow the manufacturer's instructions in the exact order.
- Follow required timing for testing.
- ☐ Identify and correct problems before reporting test results.
- Identify and report critical values in a timely manner.
- Perform or refer confirmatory or additional testing, if needed.
- ☐ Make sure patient reports are legible and reported in a timely manner.
- Make sure reports are standardized and easily distinguishable from referral laboratory test reports.
- ☐ Report patient test results only to authorized persons.
- □ Document verbal reports, followed by a written test report.
- ☐ Report public health diseases.
- □ Dispose of biohazardous waste safely.
- ☐ Participate in proficiency testing (PT).

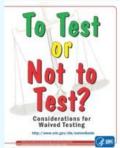


Tips, Reminders, and Resources - 11

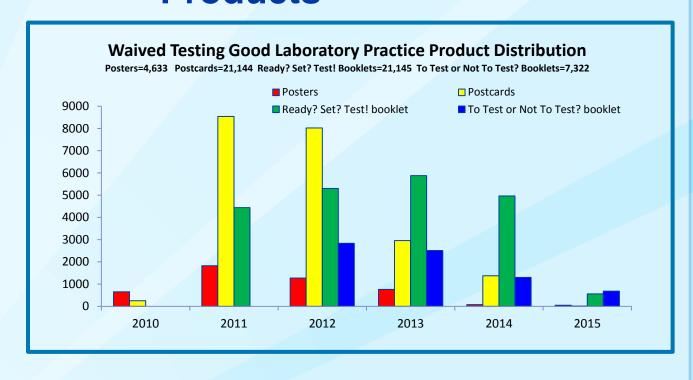
Distribution of Waived Testing Educational Products











Ready? Set? Test! Online Course Participation (Sept 2011- March 2015)

Course Registration	
Total Registered	4293
Completed	3005
In Progress	1173
Withdrawn	115

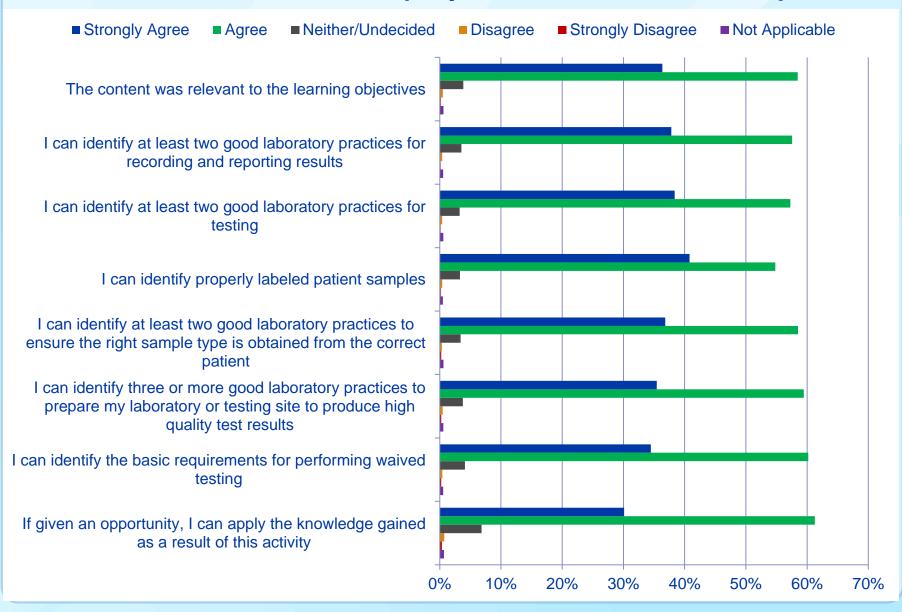
Credit Type	Total Hours Awarded
CEU/CE (0.1 hours)	127.1
CME (1.0 hours)	234
CNE Contact Hours (1.0 hours)	397
Pharmacists Contact Hours (0.1 hours)	6.3

Ready? Set? Test! Online Course

- Free training based on booklet promoting good laboratory practices for waived testing
- Includes knowledge checks and realistic waived testing scenarios to reinforce learning
- Templates, checklists, resources to save/download



Course Evaluation (Sept 2011 - March 2015)





Cytology Workload Assessment and Measure

- Purpose evaluate gynecologic cytology workload requirements for cytotechnologists using imageassisted devices
- American Society for Cytotechnology Services, Inc. – onsite observations of image-assisted screening in 8-hour workday
- Status of Time Measure Study
 - 71 Laboratory Observations of Cytotechnologists to be completed by 4/30/2015 – need 29 more volunteers
 - To participate, contact: <u>Timemeasure@asctservices.com</u>
 - More information available at:
 http://asctservices.com/cytology-workload-study/fags-time-measure-study

Workload Assessment Project Next Steps

- Partnering with schools of cytotechnology to collect student data to estimate expected workloads for newly trained employees
- CDC will provide a copy of time measure tool to each school that participates
- Students in their last month of training will participate
- CDC will collect and analyze all data
- Compare data to ASCT contracted data
- Target date to begin May 2015

Clinical Laboratory Integration into Healthcare Collaborative (CLIHC™)

CDS Tools

- Vetting new coagulation test selection algorithms
 - New anticoagulants
 - Interruption of treatment
 - Reversal of anticoagulation
 - Targeting specific conditions (VTE, DVT, PE, etc.)

Mobile applications

- CDC GA Tech Seed Grant
 - Translating coagulation algorithms to mobile app
- Evaluating Mobile Application CDS utility
 - PTT Advisor app
 - Veteran's Administration hospital physicians pilot

CLIHC™ Projects

Emory Medical Quality Improvement

 Survey Emory clinicians to better understand their laboratory test nomenclature challenges and solutions

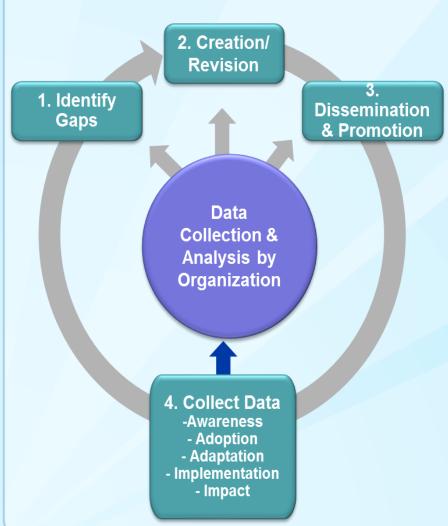
Education

 A 'Dueling Docs' scenario based skit highlighting importance of clear communication for appropriate laboratory test utilization is being developed for the CLIHC website

Partnerships

- Institute of Medicine (IOM): Diagnostic Errors in Health Care –
 Report expected 2015 with laboratory focus
- Society to Improve Diagnosis in Medicine (SIDM) Coalition to Improve Diagnosis
- CLMA's Increasing Clinical Effectiveness (ICE) coalition
- Coordinating Council on the Clinical Laboratory Workforce (CCCLW)'s Measuring Value Initiative

Laboratory Practice Guidelines (LPG) Metrics Projects



Project Goals

- Improve uptake and use of LPGs
- Identify gaps in awareness/use
- Partners develop surveys to better understand gaps and strategies to address them
- Use AGREE II tools to assess quality of representative LPGs to learn how to improve them

CDC will create a clearinghouse to promote use of LPG Metrics

- Survey design/sampling
- Focus groups
- IOM reports and related research

LPG Metrics Awardees' Projects

Clinical and Laboratory Standards Institute



- Evaluate awareness, distribution, and utility of two POC glucose monitoring (GM) LPGs for sites with (POCT 12) and without (POCT 13) laboratory support
 - LPGs chosen because both documents are relatively unknown by most of the ~85,000 testing sites

College of American Pathologists



- Immunohistochemistry (IHC) Assay Validation
 - Follow-up survey to CAP 2010 assessment
 - IHC labs to include CAP and Non-CAP PT customers
- Acute Leukemia Algorithm
 - Joint LPG with American Society of Hematology
 - 2016 Acute Leukemia Practices and Procedures Pre-Survey will provide the baseline data to measure uptake of the published LPG

LPG Metrics Awardees' Projects

- American Society for Microbiology
 - Evaluate effectiveness of four evidence-based recommendations (LMBP™)
 - Reduction of Blood Culture Contamination
 - Rapid ID of Bloodstream Infections
 - Proper Handling of Urine Specimens
 - Laboratory diagnosis of C. difficile colitis
 - Initial assessment of these LPGs by survey
 - Laboratory Response Network (LRN) to disseminate survey links
 - May establish use of LRN for surveys of sentinel clinical labs
 - Enhance public-private linkages in the states



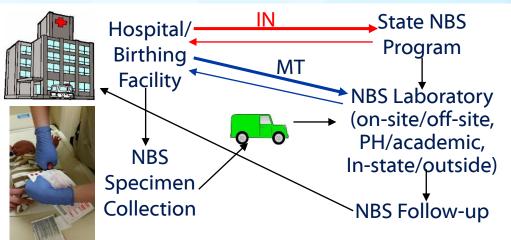
Newborn Screening (NBS) Laboratory Testing

Improve NBS - integrate birth notification with NBS specimen collection, submission, transport, process tracking to improve timeliness & quality



Current milestones:

- MT: Established 2-way messaging system between state NBS lab and birthing facility
- IN: 3 large hospital systems + 4 additional hospitals participating; birth notification/filter paper card match rates being analyzed





Laboratory Medicine Best Practices (LMBP™)

Systematic reviews in progress:

- Laboratory Test Utilization: What practices can Labs use to support and optimize appropriate Lab Test Utilization (the right test for the right patient at the right time?)
 - Practices to assess:
 - Computerized Physician Order Entry (CPOE), CDS, education, feedback, test review, lab test utilization teams, reflex testing
 - Results pending at discharge
 - Specimen rejection

LMBP™ ASM Systematic reviews collaboration

Completed

- Blood Stream Infection, Rapid Identification Methods
- Urine Specimen Handling Collection and Transport

New

- Clostridium difficile 4 focus areas:
 - 1. Highest diagnostic accuracy testing algorithms
 - 2. Increased diagnostic yield for repeat of enzyme immunoassay (EIA) or nucleic acid amplification testing
 - 3. Clinical utility of toxigenic culture and cytotoxin assay
 - 4. Clinical utility of PCR-positive and cytotoxin-/EIA-negative results

Genetic Testing Reference Materials Program (Get-RM)

Targets Characterized in Human Reference Materials for Quality Genetic Testing

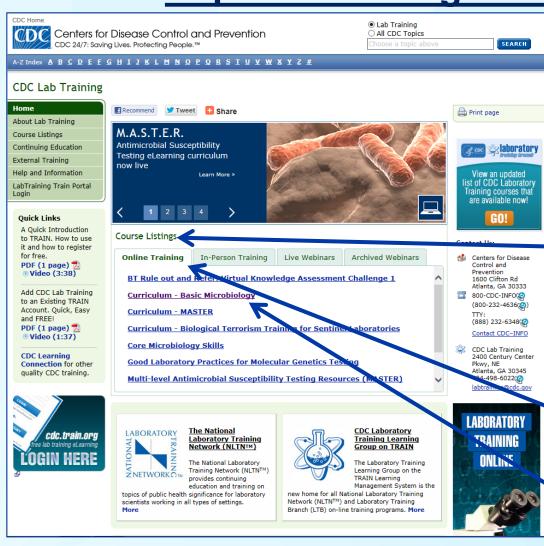


- ~700 cell lines for > 25 human genetic disorders characterized and publicly available
- New: HLA Loci and expanded pharmacogenomics haplotype characterized
 - Formed international workgroup to harmonize nomenclature and assay design

Standards for Next-Generation Sequencing in Clinical Laboratories

- CDC-facilitated national workgroups
 - "Design and Optimization of a Next-Generation Sequencing Informatics Pipeline" (in press, Nature Biotechnology)
 - A Clinical Grade Variant Template as a model for Sharing Genomic Data generated within the clinical laboratory (e.g., for external quality assurance, to outsource analysis, etc.) (manuscript in preparation)
- IOM Action Collaborative (Roundtable on Translating Genomic-based Research for Health)
 - Developing Guiding Principles for Integrating Genomic Information Into the Electronic Health Record Ecosystem

CDC Laboratory Training Website http://www.cdc.gov/labtraining/



- Access to over 50
 Online Courses
- Access to live course schedules and announcements
- Course Listings section lists all courses available from the CDC Laboratory Training Group
- Select a Course Format

 Tab
- Select a Course Topic

Highlighted Training

- The Methods of Antimicrobial Susceptibility Testing Educational Resource (MASTER)
- Basic Microbiology Blended Learning Curriculum
 - Self -study eLearning modules for didactic course content
 - Locally mentored hands-on exercises for experiential course content.
- Preparedness
 - Sentinel Laboratory "Rule out or Refer" Training
 - Sentinel Laboratory "Rule out or Refer" Training Virtual Knowledge Assessment
 - "Packaging and Shipping of Division 6.2 Materials"



For more information please contact Centers for Disease Control and Prevention

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Visit: www.cdc.gov | Contact CDC at: 1-800-CDC-INFO or www.cdc.gov/info

The findings and conclusions in this report are those of the authors and do not necessarily represent the official position of the Centers for Disease Control and Prevention.

